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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/970,049	10/02/2001	Chih-Ming Chen	300.1033US	8670
23280	7590	03/31/2005	EXAMINER	
DAVIDSON, DAVIDSON & KAPPEL, LLC 485 SEVENTH AVENUE, 14TH FLOOR NEW YORK, NY 10018			OH, SIMON J	
			ART UNIT	PAPER NUMBER
			1615	

DATE MAILED: 03/31/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

## Office Action Summary

Application No.

09/970,049

Applicant(s)

CHEN, CHIH-MING

Examiner

Simon J. Oh

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

### Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

### Status

- 1) ☒ Responsive to communication(s) filed on 26 March 2004.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

### Disposition of Claims

- 4) ☒ Claim(s) 3,6,16,17,19,20,22-25 and 27-35 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 3,6,16,17,19,20,22-25 and 27-35 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

### Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

### Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
  - ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

### Attachment(s)

- |  |   |
|--|---|
| 1) <input type="checkbox"/> Notice of References Cited (PTO-892)   | 4) <input type="checkbox"/> Interview Summary (PTO-413)<br>Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)                                   | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152)             |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)<br>Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____  |

## **DETAILED ACTION**

### ***Papers Received***

Receipt is acknowledged of the applicant's amendment and response, both received on 26 March 2004.

### ***Claim Rejections - 35 USC § 103***

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

The rejection of Claim 26 under 35 U.S.C. 103(a) as being unpatentable over the combined disclosures of Källgren, Depui *et al.*, and Eek is rendered moot with the cancellation of that claim.

The rejection of Claims 3, 6, 16, 17, 19, 20, 22-25, and 27-33 under 35 U.S.C. 103(a) as being unpatentable over the combined disclosures of Källgren, Depui *et al.*, and Eek is maintained.

Claims 34 and 35 are rejected under 35 U.S.C. 103(a) as being unpatentable over the combined disclosures of Källgren, Depui *et al.*, and Eek.

The Källgren patent teaches blister pack comprising at least a first and second row of blisters, perforated in such a way that individual blisters may be individually separated from the pack (See Abstract; Column 2, Lines 38-52; and Figures). The disclosed blister pack may be used for drugs such as omeprazole. Additionally, the blister pack is useful for packaging drugs that should be administered in combination (See Column 3, Lines 4-41).

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The Källgren patent does not explicitly teach the use of the disclosed pack with a combination of a proton pump inhibitor and a non-steroidal anti-inflammatory drug.

The Depui *et al.* patent teaches a drug combination comprising a proton pump inhibitor and a non-steroidal anti-inflammatory drug (See Abstract). Omeprazole and lansoprazole are listed as suitable proton pump inhibitors; naproxen is listed as a suitable non-steroidal anti-inflammatory drug (See Column 6, Line 1 to Column 8, Line 13). A tablet comprising lansoprazole and naproxen is disclosed (See Example 4). The use of these drugs in separate dosage forms in a combination therapy in the prior art is acknowledged in the disclosure (See Column 2, Lines 32-40). Suitable dosage ranges for each category are listed; each dosage form will preferably comprise 10 to 80 mg of the proton pump inhibitor and 10 to 800 mg of the non-steroidal anti-inflammatory drug (See Column 14, Lines 7-25)

The Eek document discloses drug packaging consisting of blister pack cards that may be assembled to form a combination pack of dosage forms, such as tablets (See Abstract; Page 1, Lines 5-12; and Figures). The scope of the disclosed invention encompasses dosage units of different drugs or different amounts of drugs within a single blister pack (See Page 5, Lines 8-14). Digital notation may be printed on the pack for the benefit of the patient. Alternatively, other notation may be printed, such as the time of day or the day of the week for the dose to be taken (See Page 7, Lines 7-11). Methods of treating disease using a combination blister pack are also disclosed (See Page 5, Lines 1-6).

It would be obvious to one of ordinary skill in the art at the time the instantly claimed invention was made to combine the disclosures of Källgren, Depui *et al.*, and Eek into the objects of the instantly claimed invention. It is the position of the examiner that one of ordinary

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skill would be motivated to combine the disclosures of Källgren, Depui *et al.*, and Eek in order to create a packaging system comprising a proton pump inhibitor in combination with a non-steroidal anti-inflammatory drug. As stated in Depui *et al.*, the administration of a non-steroidal anti-inflammatory drug in combination with a proton-pump inhibitor is known and that patient compliance is a main factor in devising a successful treatment. It is the position of the examiner that similarly, a combination dosage regimen given in a packaging system designed for that purpose, as disclosed in Källgren and Eek, will also lead to greater patient compliance. It is the position of the examiner that one of ordinary skill in the art that would recognize that the aims of the Källgren, Depui *et al.*, and Eek are similar in the area of improving patient compliance. As the disclosed invention of Källgren is not limited to any particular types of drugs to be packaged, one of ordinary skill can expect to create a drug pack comprising dosages of lansoprazole and naproxen in accordance with a combination dosage regimen with a reasonable expectation of success.

Thus, the instantly disclosed invention is *prima facie* obvious

### ***Response to Arguments***

Applicant's arguments filed 26 March 2004 have been fully considered but they are not persuasive.

In response to applicant's arguments against the references individually, one cannot show nonobviousness by attacking references individually where the rejections are based on combinations of references. See *In re Keller*, 642 F.2d 413, 208 USPQ 871 (CCPA 1981); *In re Merck & Co.*, 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986).

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Furthermore, the present amendments to the claims are primarily directed to indicia that are to be included in the instantly claimed drug packaging system. It has been recently established that instructions included on packaging that are not functionally related to a product, but merely show a use for an existing product, do not constitute a patentable invention. *In re Ngai and Lin* (U.S. Court of Appeals for the Fed. Cir. 03-1524, decided March 8, 2004).

### ***Conclusion***

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire **THREE MONTHS** from the mailing date of this action. In the event a first reply is filed within **TWO MONTHS** of the mailing date of this final action and the advisory action is not mailed until after the end of the **THREE-MONTH** shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than **SIX MONTHS** from the date of this final action.

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*Correspondence*

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Simon J. Oh whose telephone number is (571) 272-0599. The examiner can normally be reached on M-F 8:30 am to 5:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Thurman K. Page can be reached on (571) 272-0602. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Simon J. Oh  
Examiner  
Art Unit 1615

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